Amendments to claims

This listing of claims will replace all prior versions and listing of claims in the application.

Please cancel claim 1 without prejudice or disclaimer and add new claims 39 through 83 as shown.

1-38. (canceled).

39. (new): An anhydrous composition formulated for topical delivery comprising:

(a) an alcohol selected from the group consisting of ethanol, isopropyl alcohol and isobutanol,

(b) propylene glycol,

(c) polyethylene glycol,

(d) glycerin, and

(e) ketoconazole,

wherein the composition is formulated as an anhydrous gel and does not contain a retinoid.

40. (new): The composition of claim 39 wherein the ketoconazole is solubilized.

41. (new): The composition of claim 39 wherein the amount of the ketoconazole is about 0.5 to about 3 percent by weight

42. (new): The composition of claim 39 wherein the amount of the ketoconazole is about 2.0 percent by weight.

43. (new): The composition of claim 39 wherein the alcohol is ethanol.

44. (new): The composition of claim 39 wherein the amount of the polyethylene glycol is about 10 to about 80 percent by weight.

45. (new): The composition of claim 44 wherein the amount of the polyethylene glycol is about 20 percent by weight.

46. (new): The composition of claim 39 wherein the amount of the propylene glycol is about 1.0 to about 50 percent by weight.

47. (new): The composition of claim 46 wherein the amount of the propylene glycol is about 20 percent by weight.

48. (new): The composition of claim 39 wherein the amount of the glycerin is about 10 to about 80 percent by weight.

49. (new): The composition of claim 48 wherein the amount of the glycerin is about 20 percent by weight.

50. (new): The composition of claim 39 wherein the composition further comprises an emollient.

51. (new): The composition of claim 50 wherein the amount of the emollient is between 0 and about 10 percent by weight.

52. (new): The composition of claim 50 or claim 51, wherein the emollient is PPG-15 stearyl ether.

53. (new): The composition of claim 52 wherein the amount of the PPG-15 stearyl ether is between 0 and about 2 percent by weight.

54. (new): The composition of claim 53 wherein the amount of the PPG-15 stearyl ether is about 2 percent by weight.

55. (new): The composition of claim 39 wherein the composition further comprises a viscosifier.

56. (new): The composition of claim 55 wherein the amount of the viscosifier is between 0 and about 5 percent by weight.

57. (new): The composition of claim 55 or claim 56 wherein the viscosifier is hydroxypropyl cellulose.

58. (new): The composition of claim 57 wherein the amount of hydroxypropyl cellulose is about 1.5 to about 2.0 percent by weight.

59. (new): The composition of claim 39 wherein the composition further comprises a pH adjuster.

60. (new): The composition of claim 59 wherein the amount of the pH adjuster is between 0 and about 2 percent by weight.

61. (new): The composition of claim 59 or claim 60 wherein the pH adjuster is selected from the group consisting of ascorbic acid, citric acid and combinations thereof.

62. (new): The composition of claim 61 wherein the amount of the ascorbic acid is between 0 and about 0.3 percent by weight.

63. (new): The composition of claim 61 wherein the amount of the citric acid is between 0 and about 0.5 percent by weight.

64. (new): The composition of claim 62 wherein the amount of the ascorbic acid is about 0.3 percent by weight.

65. (new): The composition of claim 63 wherein the amount of the citric acid is about 0.1 percent by weight.

66. (new): The composition of claim 39 wherein the composition further comprises an antioxidant.

67. (new): The composition of claim 66 wherein the amount of the antioxidant is between 0 and about 2 percent by weight.

68. (new): The composition of claim 66 or claim 67 wherein the antioxidant is selected from the group consisting of ascorbic acid, butylated hydroxytoluene and combinations thereof.

69. (new): The composition of claim 68 wherein the amount of the butylated hydroxytoluene is between 0 and about 0.1 percent by weight.

70. (new): The composition of claim 69 wherein the amount of the butylated hydroxytoluene is about 0.1 percent by weight.

71. (new): The composition of claim 39 wherein the composition further comprises one or more colorants.

72. (new): An anhydrous composition formulated for topical delivery comprising:

(a) propylene glycol,

(b) polyethylene glycol,

(c) glycerin,

(d) ethanol,

(e) ketoconazole,

(f) PPG-15 stearyl ether,

(g) hydroxypropyl cellulose,

(h) ascorbic acid,

(i) butylated hydroxytoluene, and

(j) citric acid,

wherein the composition is formulated as an anhydrous gel and does not contain a retinoid.

73. (new): An anhydrous composition formulated for topical delivery consisting essentially of:

(a) propylene glycol,

(b) polyethylene glycol,

(c) glycerin,

(d) ethanol,

(e) ketoconazole,

(f) PPG-15 stearyl ether,

(g) hydroxypropyl cellulose,

(h) ascorbic acid,

(i) butylated hydroxytoluene, and

(j) citric acid,

wherein the composition is formulated as an anhydrous gel.

74. (new): A method of delivering a composition of any one of claims 39, 72 and 73 for the treatment of skin fungal disorders to a recipient in need of such treatment comprising topically administering the composition to the recipient.

75. (new): The method of claim 74 wherein the recipient is a human.

76. (new): The method of claim 75 wherein the human is suffering from seborrheic dermatitis.

77. (new): A method of treating skin fungal disorders comprising topically administering the composition of any one of claims 39, 72 and 73 to a recipient in need of such treatment.

78. (new): The method of claim 77 wherein the recipient is a human.

79. (new): The method of claim 78 wherein the skin fungal disorders are associated with *T. rubrum* or *P. ovale*.

80. (new): The method of claim 79, wherein the skin fungal disorders are selected from the group consisting of tinea corporis, tinea cruria, tinea pedis and seborrheic dermatitis.

81. (new): A method of treating seborrheic dermatitis comprising topically administering the composition of any one of claims 39, 72 and 73 to a recipient in need of such treatment.

82. (new): The method of claim 81 wherein the recipient is a human.

83. (new): The method of claim 74 wherein the skin fungal disorders are associated with *T. rubrum* or *P. ovale*.